



Chirality in microbial biofilms is mediated by close interactions between the cell surface and the substratum

Jauffred, Liselotte; Vejborg, Rebecca Munk; Korolev, Kirill S.; Brown, Stanley; Oddershede, Lene B.

Published in:
I S M E Journal

DOI:
[10.1038/ismej.2017.19](https://doi.org/10.1038/ismej.2017.19)

Publication date:
2017

Document version
Publisher's PDF, also known as Version of record

Citation for published version (APA):
Jauffred, L., Vejborg, R. M., Korolev, K. S., Brown, S., & Oddershede, L. B. (2017). Chirality in microbial biofilms is mediated by close interactions between the cell surface and the substratum. *I S M E Journal*, 11(7), 1688-1701. <https://doi.org/10.1038/ismej.2017.19>

ORIGINAL ARTICLE

Chirality in microbial biofilms is mediated by close interactions between the cell surface and the substratum

Liselotte Jauffred¹, Rebecca Munk Vejborg², Kirill S Korolev³, Stanley Brown¹ and Lene B Oddershede¹

¹The Niels Bohr Institute, University of Copenhagen, Copenhagen, Denmark; ²Costerton Biofilm Center, University of Copenhagen, Copenhagen, Denmark and ³Department of Physics and Graduate Program in Bioinformatics, Boston University, Boston, USA

From microbial biofilms to human migrations, spatial competition is central to the evolutionary history of many species. The boundary between expanding populations is the focal point of competition for space and resources and is of particular interest in ecology. For all *Escherichia coli* strains studied here, these boundaries move in a counterclockwise direction even when the competing strains have the same fitness. We find that chiral growth of bacterial colonies is strongly suppressed by the expression of extracellular features such as adhesive structures and pili. Experiments with other microbial species show that chiral growth is found in other bacteria and exclude cell wall biosynthesis and anisotropic shape as the primary causes of chirality. Instead, intimate contact with the substratum is necessary for chirality. Our results demonstrate that through a handful of surface molecules cells can fundamentally reorganize their migration patterns, which might affect intra- and interspecific competitions through colony morphology or other mechanisms.

The ISME Journal (2017) 11, 1688–1701; doi:10.1038/ismej.2017.19; published online 14 April 2017

Introduction

From the structure of amino acids to the shape of our galaxy, chirality is ubiquitous in nature and has fascinated scientists for centuries. The accepted mechanism for the emergence of left–right asymmetry is a fluctuation producing a chiral state followed by positive feedback favoring homochirality (Frank, 1953; Gayathri and Rao, 2005; Saito *et al.*, 2007). More recent work further demonstrates that intrinsic noise in autocatalytic reactions is also sufficient to produce and stabilize left–right asymmetry (Jafarpour *et al.*, 2015). While these physical mechanisms might explain chirality in the inanimate world, chirality in living systems could be a product of natural selection.

Although the ecological and evolutionary role of chirality is largely unexplored, there are several examples that point to the possibility that chirality could be advantageous. Extensive work with *Paenibacillus* showed that this microbe switches from chiral to non-chiral phenotypes in response to changing environmental conditions (Ben-Jacob *et al.*, 2000). More recently, Wan *et al.* (2011) found

that cancer cells are chiral and have the opposite chirality of the normal tissue from the same patient. In both cases, molecular changes propagate to population-level scales and alter the ecology and evolution of the species. The details of these processes are poorly understood. These striking examples motivated us to develop a molecular understanding of chirality in a simple model system of an *Escherichia coli* colony.

In the lab, *E. coli* typically grows as a compact circular colony when inoculated on an agar plate; yet internal dynamics of those colonies are often chiral. These dynamics can be revealed by fluorescently labeling subpopulations of growing cells. Strong genetic drift at the growing edge of the colony promotes local loss of genetic diversity and results in spatial de-mixing of the subpopulations producing a characteristic pattern of flaring sectors on long timescales (>30 generations) (Kreft, 2004; Hallatschek *et al.*, 2007; Ali and Grosskinsky, 2010; Hallatschek and Nelson, 2010; Korolev *et al.*, 2010). Without chirality, the boundaries between the sectors extend radially, but, in chiral species, boundaries appear as spirals emerging from the center of the colony (Korolev *et al.*, 2011).

While the existence of chirality in bacterial colony growth has been established, there are still open and interesting questions regarding the origin and evolutionary importance of sector boundary chirality:

Correspondence: LB Oddershede, Niels Bohr Institute, University of Copenhagen, Blegdamsvej 17, Copenhagen 2100, Denmark. E-mail: oddershede@nbi.dk

Received 3 June 2016; revised 22 December 2016; accepted 18 January 2017; published online 14 April 2017

Table 1 Description of the strains and plasmids used in the study

Name, relevant characteristics		Reference
<i>E. coli</i> strains		
DH5 α	LuxS supE44 Δ lacU169, (ϕ 80 lacZ Δ M15) hsdR17, recA1, endA1, gyrA96, thi-1, relA1	Invitrogen
DH5 α -H	LuxS supE44 Δ lacU169, (ϕ 80 lacZ Δ M15) hsdR17, recA1, endA1, gyrA96, thi-1, relA1	Hallatschek <i>et al.</i> (2007)
MS613	MG1655 K-12 reference (F-lambda- ilvG- rfb-50 rph-1) strain	Bachmann (1996)
MS427	MG1655 Δ flu	Reisner <i>et al.</i> (2003)
MS428	MG1655 Δ fim	Kjærgaard <i>et al.</i> (2000)
MS528	MG1655 Δ flu, Δ fim	Kjærgaard <i>et al.</i> (2000)
RMV340	MG1655cps::tet	Beloin <i>et al.</i> (2006)
RMV359	MG1655fliC::kan	This study
RMV612	MG1655csgAB::kan	This study
<i>S. pombe</i> strains		
PG3732	h+ ura4D::mCherry	G Thon
PG3733	h+ ura4D::YFP	G Thon
<i>B. subtilis</i> strains		
168phymKATE	eps- : epsG	van Gestel <i>et al.</i> (2014)
168phymGFP	eps- : epsG	van Gestel <i>et al.</i> (2014)
Plasmids		
pVS130	eCFP	V Sourjik
pVS132	YFP	V Sourjik
pKD4	ApKm, kan cassette template	Datsenko and Wanner (2000)
pKD46	Ap, λ Red recombinase	Datsenko and Wanner (2000)
pCP20	ApCm, FLP recombinase expression	Datsenko and Wanner (2000)
pACYC184	Cloning vector, Cam ^R	Chang and Cohen (1978)
pHHA13	Wild-type fim gene cluster in pACYC184	H Hasman

Abbreviations: CFP, cyan fluorescent protein; YFP, yellow fluorescent protein.

(i) why and when is chirality beneficial, (ii) how does microscopic chirality translate into a macroscopic chirality and (iii) what are the molecular origins of the chirality? The first question is of great interest and, while our work does not address this question directly, it provides important foundation for future studies. The second question has been largely resolved for bacteria via thorough mathematical modeling and experimentation (Ben-Jacob *et al.*, 2000; Xue *et al.*, 2011). Here, we primarily focus on the last question and uncover the key molecular players shaping chirality of bacterial growth.

Already in 1848, Louis Pasteur related macroscopic chirality of a crystal to a microscopic symmetry breaking of constitutive molecules (Flack, 2009). Similar mechanisms operate in biological systems; for example, *Lymnaea stagnalis* snail shell chirality could be reversed by altering asymmetry in the blastomere (Kuroda *et al.*, 2009). Although the general principle that macroscopic chirality originates at cellular or subcellular level is well established, a detailed molecular and biophysical understanding of these processes is still lacking for all but a handful of systems. Perhaps the best-studied example of chiral behavior in bacteria is that of *E. coli*, which turns clockwise while swimming near surfaces due to the chiral motion of its flagella (DiLuzio *et al.*, 2005).

To uncover the molecular mechanisms of chiral growth, we studied several microbial species and *E. coli* mutants lacking certain surface structures under varying environmental conditions. Our results ruled out shape and the direction of cell wall biosynthesis as the primary drivers of chiral growth. Instead, the data suggest that chirality relies on intimate contact between the substratum and the *E. coli* cell surface. This contact can be masked by surface structures; thus, the deletion of surface factors progressively increases *E. coli* chirality compared with wild type. Not only the cell surface, but also the substratum itself is important for the interaction mediating chirality. We found maximal chiral behavior at intermediate agar concentration (1–1.5%) and a significant dependence on substrate composition. Collectively, our results reveal how molecular mechanisms control spatial patterning during colony growth and suggest new ecological roles for several proteins expressed on cell surfaces.

Materials and methods

Cell culture

The strains used in this study are listed in Table 1. *E. coli* strains were grown at 37 °C in Luria-Bertani (LB) broth (1% tryptone, 0.5% NaCl, 0.5% yeast extract) or on LB agar (1.5% agar) supplemented with the appropriate antibiotics, unless otherwise

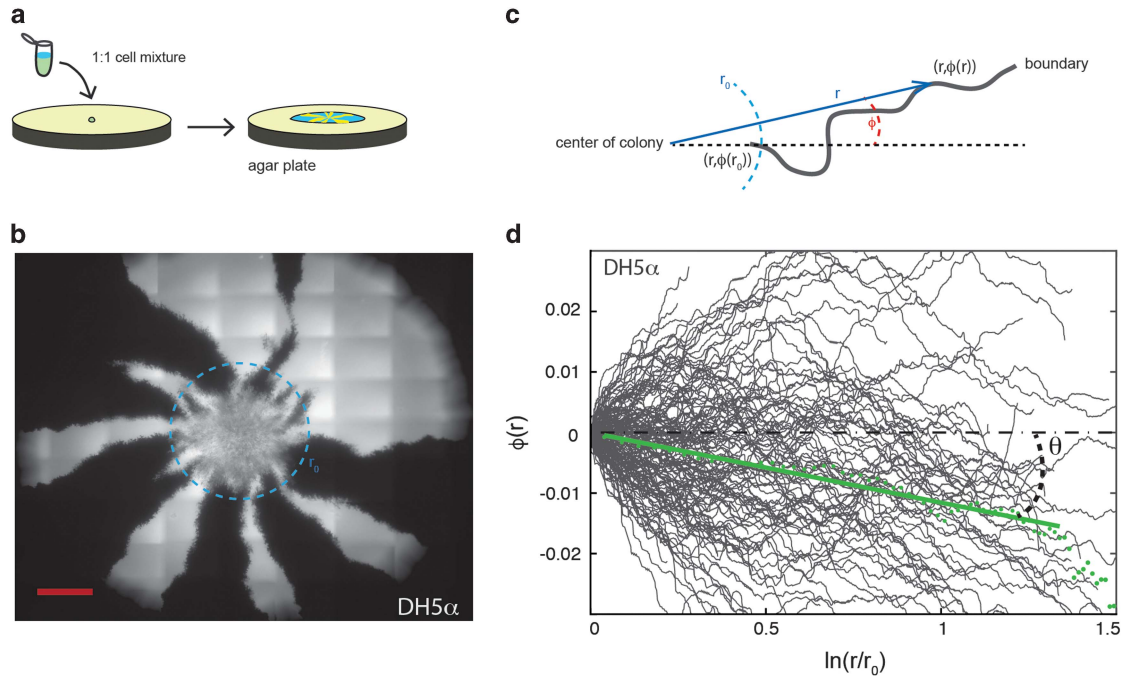


Figure 1 Competition experiments between neutral *E. coli* strains expressing different colors, either yellow fluorescent protein (YFP) (yellow) or cyan fluorescent protein (CFP) (cyan). (a) Sketch of the experiment where the two species are mixed 1:1, diluted 1/10, and inoculated on an agar plate where the borders between competing expanding strains are visible. (b) An example of an expanding *E. coli* DH5α colony is shown from the agar side; the cells transformed with CFP appear white and those with YFP black in this image. The radius of the light-blue dashed circle equals r_0 . The scale bar corresponds to 1 mm. (c) Illustration of the parameterization, $(r, \phi(r))$, of a point on the sector boundary where r is the length of the radial vector (blue) from the center of the colony and $\phi(r)$ is the angle (denoted with orange) between the direction of the sector boundary (dashed orange line) and the radial vector. (d) Parameterization, $\phi(r)$, of individual *E. coli* DH5α sector boundaries ($N=146$) plotted on semilog scales. The green dots show the average boundary, $\langle \phi(r) \rangle$, and the full green line shows the best linear fit to the average boundary. As clear from Equation (1), the slope of the fit is the dimensionless measure of chirality: $v_{\perp}/v_{\parallel} = \tan(\theta)$. Note that, although the behavior of individual boundaries appear erratic due to random wandering of the boundaries, the average over 146 colonies shows a clear negative slope due to chirality. The values of v_{\perp}/v_{\parallel} and θ for all strains investigated are given in Table 2.

stated. *Bacillus subtilis* and *Schizosaccharomyces pombe* were grown at 30 °C in LB and YPD broth supplemented with uracil, respectively. For competition experiments, differently labeled *E. coli* K-12 strains were obtained by transformation with plasmids pVS132 and pVS130 expressing yellow fluorescent protein or cyan fluorescent protein (Hallatschek *et al.*, 2007), respectively. Antibiotics were used at the following concentrations: kanamycin (25 $\mu\text{g ml}^{-1}$), ampicillin (*E. coli*: 100 $\mu\text{g ml}^{-1}$, *B. subtilis*: 10 $\mu\text{g ml}^{-1}$). For the MS528+pHHA13 strain, ampicillin (100 $\mu\text{g ml}^{-1}$) and chloramphenicol (10 $\mu\text{g ml}^{-1}$) were added to the agar and medium. For cells grown on trypsin plates we added 0.025% trypsin w/v (Sigma Aldrich, Darmstadt, Germany) to the LB agar together with Isopropyl β -D-1-thiogalactopyranoside (IPTG) and ampicillin.

Competition experiments

The strains were grown as separate cell cultures. Overnight cultures were mixed 1:1 and 1 μl droplets were inoculated on plates with ampicillin (100 $\mu\text{g ml}^{-1}$) and IPTG (0.5 mM for induction of fluorophores). The seeding of a droplet on an agar plate is illustrated in Figure 1a. *B. subtilis* was grown on plates with ampicillin (10 $\mu\text{g ml}^{-1}$) with or

without IPTG (0.5 mM for exo-polysaccharide induction (van Gestel *et al.*, 2014)). We also did line inoculations by dipping the side of a cover slip in the cell culture and gently placing it on the agar plate afterwards. Plates with partly absorbed inoculation droplets were incubated at 37 °C for up to 11 days or until the circular colony had obtained a diameter of at least 1.5–2 cm. The cell density used for the inoculation influenced the patchiness of the homeland (Hallatschek *et al.*, 2007) and the number of resulting sectors (van Gestel *et al.*, 2014), but had no apparent effect on the sectoring pattern as also reported (Hallatschek *et al.*, 2007).

Recombinant DNA techniques

Specific mutants (MG1655/*fliC* and MG1655/*csgAB*) of *E. coli* MG1655 were constructed using the λ -red recombinase-based gene inactivation system as described (Datsenko and Wanner, 2000; Yu *et al.*, 2000). Following amplification of the kanamycin cassette from pKD4, the PCR products were transformed into MG1655 carrying the helper plasmid pKD46 and the transformations were plated on selective plates (50 $\mu\text{g ml}^{-1}$ kanamycin; primer sequences are given in supplementary Table S1). Deletion mutants were verified by PCR using the test

primers given in supplementary Table S1. The kanamycin-resistance cassette was subsequently removed by use of the Flippase-encoding plasmid, pCP20 (Datsenko and Wanner, 2000). For complementation experiments, the pHHA13 plasmid was transformed into the MS528 strain.

Trypsin assay

To test whether trypsin modulates the cell surface or the substratum, autoclaved agar and water were mixed into freshly filter-sterilized LB broth with or without trypsin (0.025%) and left overnight at room temperature. On the following day, the media were

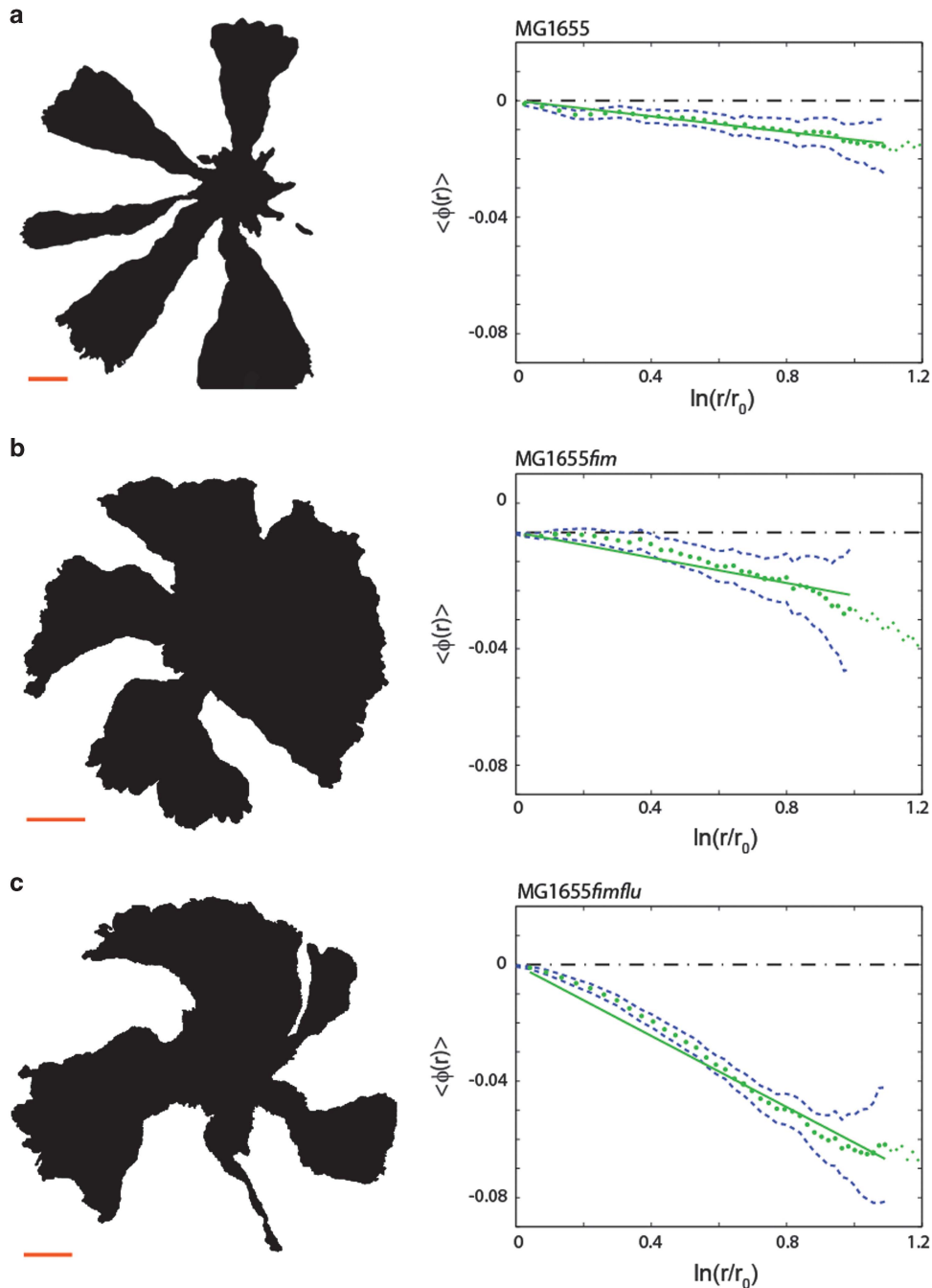


Figure 2 The sector boundaries show different degrees of chirality depending on expression of surface structures. The left panels show examples of intensity thresholded images (displayed from the agar side) of different *E. coli* MG1655 mutants in competition experiments; the scale bars are 1 mm. The right panels show the average boundary angle, $\langle \phi(r) \rangle$ (green dotted line), a linear fit (full green line) and one standard error of the mean (blue punctuated line). (a) Results from competition experiments with the mother strain, MG1655; this colony expansion is only slightly chiral ($N=369$). (b) Competition experiments with the MG1655*fim* strain lacking pili ($N=831$). This strain exhibits more chiral borders than MG1655. (c) Competition experiments with the double deleted MG1655*fimflu* strain lacking both Antigen 43 and pili ($N=636$). This double deletion causes even more pronounced chirality compared with the single deletion mutants.

autoclaved and IPTG and ampicillin were added. As trypsin under some conditions is heat-resistant (Edie, 1914), trypsin-inhibitor AEBSF (0.02%) was added to half of the medium before pouring plates (1.2% agar). A third medium was prepared from fresh powders and no trypsin.

Substratum substitutions

To identify which of the ingredients of LB agar are responsible for chiral growth, we substituted each ingredient one by one (while always maintaining IPTG and ampicillin). Yeast extract was replaced by $1 \mu\text{g ml}^{-1}$ vitamin B₁, 1 mM MgCl₂ and 10 mM sodium phosphate. Tryptone and NaCl were replaced by 1% decolorized casamino acids (which also contained NaCl). Agar (1.2%) was replaced by 1.2% agarose. Notably, these experiments were done with a viscosity of 1.2% agar/agarose.

Fluorescence microscopy

The colonies were imaged with an inverted Nikon Eclipse Ti fluorescent microscope (Nikon, Tokyo, Japan) using a $\times 4$ air objective and an Andor Neo camera (Andor, Belfast, UK). Emission from yellow fluorescent protein/cyan fluorescent protein was collected at 535/485 nm (30 nm bandwidth) upon excitation at 500/405 nm (20 nm bandwidth). The agar plates were placed upside down directly on the microscope and images were obtained in an array of 12×12 sub-images. Individual sub-images were automatically stitched after acquisition with a 15% overlap to image the entire colony, which typically had a diameter of 2–4 cm.

Transmission electron microscopy

Bacteria were harvested from semisolid agar plates and analyzed with respect to their cell morphology and the possible presence of flagella using transmission electron microscopy. The cells were adsorbed onto carbon-coated grids and negatively stained with 2% phosphotungsten acid. The samples were then visualized with a Philips CM 100 TEM (Philips, Eindhoven, The Netherlands) equipped with a OSIS Veleta digital slow scan $2 \text{ k} \times 2 \text{ k}$ CCD camera. Digital images were recorded using the ITEM software package (Irvine, CA, USA).

Tracking of sector boundaries

In fluorescent microscopy images of bacterial colonies, the sectors were visible as shown in Figure 1b. The images were converted to black and white by an automated Matlab thresholding procedure (see, for example, left part of Figure 2). Next, we identified the center of the colony and the radius, r_0 , at which clear sector boundaries first appear, by fitting a circle to the colony rim using the Taubin method in Matlab. A circle with radius r_0 is marked on the image in Figure 1b, the strains are mixed inside this

region, but sectors can be clearly identified outside r_0 , this radius being limited by nutrients (Mitri *et al.*, 2015). The mixed region was typically larger than the original inoculated droplet and r_0 should be subscript. Then, each colony boundary was identified by edge detection and each pixel on a boundary was assigned a radius (distance to center), r , and an angle, φ . The latter denotes the angle between the boundary and the radial vector, see definitions of r and φ in Figure 1c. Thus, the individual boundaries were parametrized by custom-made Matlab procedures and the average boundary for each data set, $\langle \varphi(r) \rangle$, was found by binning and averaging over all boundaries. The sector boundaries were found to twist as an equiangular, that is, logarithmic, spiral (Huntley, 1970), and each average boundary, $\langle \varphi(r) \rangle$, was therefore least-squared fitted to Equation (1) (Korolev *et al.*, 2011), which describes the evolution of an equiangular spiral:

$$\langle \varphi(r) \rangle = \varphi(r_0) + \frac{v_{\perp}}{v_{\parallel}} \ln\left(\frac{r}{r_0}\right). \quad (1)$$

Here, v_{\parallel} is the velocity in the radial direction, and v_{\perp} the velocity in the direction perpendicular to the expanding frontier. In the following, we only consider the average change in boundary angle $\varphi(r) - \varphi(r_0)$ as a function of the radius r , but denote it as $\langle \varphi(r) \rangle$ for simplicity. This can be interpreted as simply choosing the reference direction along $\langle \varphi(r_0) \rangle$. For the DH5 α strain, the spatial pattern of rotating sector boundaries is shown in Figure 1b and its analysis in terms of Equation (1) is shown in Figure 1d where individual boundaries are plotted as full gray lines and the average boundary is given by green dots. The data are indeed consistent with a linear dependence of a semilog plot as predicted by Equation (1). The slope of the line equals v_{\perp}/v_{\parallel} , and thus quantifies chirality. In the following, we use a more convenient measure of chirality θ defined as $v_{\perp}/v_{\parallel} = \tan(\theta)$ (see definition of θ in Figure 1d). θ denotes the angle between the outward expansion of the colony and the direction of boundary motion. We only observed left-handed (sinistral) motion of sector boundaries when seen from the bottom of the colony (the agar side), so all chiral angles reported are negative. The larger the absolute value of θ , the more chiral the colonies.

Results

Observation of chiral boundaries between competing strains

In this study, we used pairs of isogenic *E. coli* K-12 strains expressing two different fluorescent proteins (yellow fluorescent protein/cyan fluorescent protein) and the strains with the two labels have similar (neutral) fitness (Hallatschek and Nelson, 2010; Korolev *et al.*, 2011). For competition experiments, a mixture of the two neutral strains was inoculated on an agar surface as illustrated in Figure 1a and

colony expansion was quantified by fluorescence microscopy. The growing colonies exhibit a striking segregation of the two populations (cyan fluorescent protein and yellow fluorescent protein) over time in accordance with previous observations. The boundaries, resulting from population segregation, were identified and the chirality angle, θ , determined. The example shown in Figures 1b and d is from an *E. coli* DH5 α colony. The dynamics of segregation was restricted to the edge of the colony, whereas, except for a gradual thickening, the interior distribution of cells was frozen as previously reported (Hallatschek and Nelson, 2010; Korolev *et al.*, 2011). Our analysis shows that DH5 α colonies have chiral sector boundaries with $\theta = -6.1^\circ$ (s.e.m. = 0.05° , $N = 146$), where N is the number of boundaries over all investigated colonies. This compares well to the interval (from -5° to -7°) reported for the DH5 α -H strain in Hallatschek *et al.* (2007). The same strain was later shown to have a chirality angle, θ , within the interval from -13° to -17° (Korolev and Nelson, 2011), which is reasonably similar to the value obtained in the current study for the DH5 α -H strain (all θ -values are given in Table 2). This demonstrates that even between similar strains, or strains with the same original ancestry, differences in pedigree or growth conditions might affect the chirality angle.

Using different strains, we did both line inoculations (see Supplementary Figure S1) and circular inoculations (as shown in Figure 1) and observed that the sectoring occurs in both types of assays with a similar chirality of the boundaries. Hence, chirality is not linked to radial expansions of the colony front *per se* but is a more general feature of the species.

Systematic removal of extracellular structures

It is a possibility that the sector boundaries observed in bacterial colonies are related to the composition of the cell wall or the extracellular matrix. Obvious determinants could be bacterial cell adhesive surface structures such as exo-polysaccharides, fimbriae, flagella and extracellular DNA, all of

which have previously been shown to play a role in cell-surface and cell-cell interactions (see O'Toole *et al.*, 2000 for a review on biofilm formation). Based on this hypothesis, we examined the chirality of expanding colonies of a collection of *E. coli* strains with specific deletions in genes encoding various bacterial surface structures. Information on all strains and relevant plasmids used is listed in Table 1 and Figure 3 shows transmission electron microscopy images of a MG1655 cell as well as a sketch illustrating the relevant surface structures, that is, flagella, pili, colanic acid, curli fimbriae and antigen 43.

To examine the influence of such structures, a systematic examination of these cell surface components was undertaken, using the parental strain MG1655 (Bachmann, 1996). The sectoring during expansion of MG1655 is shown in Figure 2a. As visible both in the (left) image and in the (right) plot of the average boundary, MG1655 is only slightly chiral with $\theta = -0.8^\circ$ (s.e.m. = 0.003° , $N = 369$) (the values of θ for all strains are listed in Table 2). Thus, *E. coli* MG1655 (Figure 3a) displays significantly less chirality than DH5 α (Figure 1b). As DH5 α and MG1655 both are K-12 strains of *E. coli*, we find that even among closely related strains the chirality can vary substantially.

We then examined the effect of deleting different adhesion proteins: We first tested MG1655/*flu*, which carries a deletion in the gene encoding the auto-transporter protein antigen 43, a *handshake* protein which is important for cell-cell aggregation and cellular chain formation (Diderichsen, 1980; Vejborg and Klemm, 2009). Antigen 43 appears in the outer membrane of various bacterial species, including *E. coli*, and come in copy numbers up to 50 000 per cell (Owen *et al.*, 1987). For MG1655/*flu* $\theta = -0.4^\circ$ (s.e.m. = 0.05° , $N = 192$), hence, deletion of the antigen 43 gene, *flu*, lowered chirality slightly, but not significantly, with respect to the parental strain MG1655. We then tested MG1655/*fim*, a strain that does not express pili (also known as type I fimbriae) which are important for cell-cell and cell-surface adhesion. Figure 2b shows that MG1655/*fim* $\theta = -1.4^\circ$ (s.e.m. = 0.01° , $N = 831$), which is significantly more chiral than MG1655/*flu* and MG1655. Chirality was even more pronounced in the double mutant, MG1655/*flufim*; the expansion of this mutant is shown in Figure 2c and it had $\theta = -3.5^\circ$ (s.e.m. = 0.008° , $N = 636$). Hence, the chirality of the double mutant was significantly larger than a simple superposition of the chirality angles of the two individual deletion mutants. Complementing the double deletion mutant with a *fim*-encoding plasmid (MG1655/*flufim* + pHHA13) reduced the chirality as expected, although not to parental level, thus suggesting that the phenotype is multifactorial. Given the inverse regulation of pili and antigen 43 (Hasman *et al.*, 1999, 2000), the slight loss of chirality for MG1655/*flu* could be a secondary effect, for example, the *flu* mutant could be overexpressing *fim*.

Table 2 Chirality of the different strains

Strain	Name	N	$\theta \pm \text{s.e.m.}$
DH5 α	—	146	$-6.1^\circ \pm 0.05^\circ$
DH5 α -H	—	183	$-10.2^\circ \pm 0.05^\circ$
MS613	MG1655	369	$-0.8^\circ \pm 0.003^\circ$
MS427	MG1655/ <i>flu</i>	192	$-0.4^\circ \pm 0.005^\circ$
MS428	MG1655/ <i>fim</i>	831	$-1.4^\circ \pm 0.01^\circ$
MS528	MG1655/ <i>fimflu</i>	636	$-3.5^\circ \pm 0.008^\circ$
RMV340	MG1655/ <i>cps</i>	989	$-1.3^\circ \pm 0.008^\circ$
RMV612	MG1655/ <i>csgAB</i>	646	$-1.3^\circ \pm 0.009^\circ$
RMV359	MG1655/ <i>fliC</i>	542	$-0.4^\circ \pm 0.005^\circ$
MS528Complemented	MG1655/ <i>fimflu</i> + pHHA13	50	$-1.8^\circ \pm 0.05^\circ$
MS528trypsin	MG1655/ <i>fimflu</i> + trypsin	1952	$-1.0^\circ \pm 0.005^\circ$
<i>B. subtilis</i> 168	<i>B. subtilis</i>	286	$-1.1^\circ \pm 0.03^\circ$

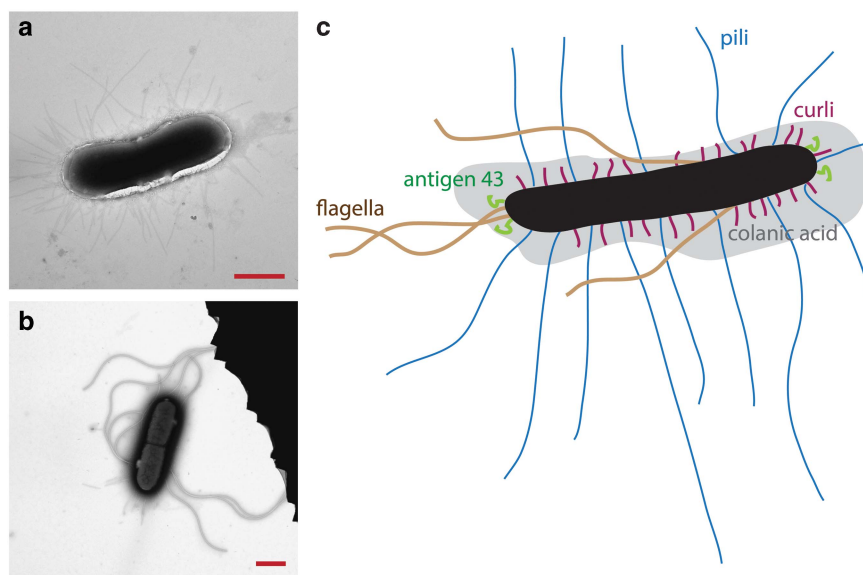


Figure 3 Electron microscopy images as well as a sketch of the *E. coli* and the surface structures investigated. (a, b) Transmission electron microscopy images of MG1655. The scale bar is 1 μm . (c) A sketch of *E. coli* with its extracellular constituents: Antigen 43, pili, curli fimbriae, colanic acid and flagella are explicitly drawn (not to scale). The body of the bacteria is shown in black.

As may be apparent from the results with MG1655*flu*, MG1655*fim* and MG1655*fimflu*, the measure of chirality used here, θ , is not additive. Given that θ is small, one might expect that Taylor's approximation holds and that the contributions from independent mutations should simply add up. However, the deviation from additivity observed here suggests that *flu* and *fim* mutations have epistatic interactions, possibly due to induced changes in gene expression, which is in line with the fact that many of the bacterial surface structures display complex regulation (Saldaña *et al.*, 2009).

Colanic acid is an exo-polysaccharide important for biofilm formation while curli fimbriae are important for cell–cell and cell–surface adhesion (Vidal *et al.*, 1998; Prigent-Combaret *et al.*, 1999, 2000; Danese *et al.*, 2000; Oh *et al.*, 2016). Both of these are linked to biofilm formation, although predominantly at ambient temperature (Prigent-Combaret *et al.*, 2000; Gualdi *et al.*, 2008). To examine the effect of the biofilm matrix structures, mutants carrying deletions in genes encoding colanic acid, MG1655*cps*, and curli fimbriae, MG1655*cgsAB*, were tested. MG1655*cps* and MG1655*cgsAB* both exhibited an increased chirality in comparison with MG1655.

We also examined the effect of flagella expression. For swimming motility in *E. coli*, a motor rotates the peritrichous flagella controlling the formation of a corkscrew-shaped bundle. By rotating the flagella counterclockwise or clockwise, and by bundling or de-bundling the flagella *E. coli* performs its characteristic run-tumble-run (Wolfe and Berg, 1989) and is able to move rapidly in response to chemical gradients. This is also the case for (chemotactic) *E. coli* cells on semisolid agar (0.20–0.35%) (Wolfe and Berg, 1989). The exact function of the flagella on solid agar, as used for this study (1.5%), is still

unknown although recent studies suggest that they may have a structural role in biofilm formation. The sector boundaries in colony formation of the flagella mutant, MG1655*fliC*, had a slightly, but not significantly, lower degree of chirality than the parental strain, $\theta = -0.4^\circ$ (s.e.m. = 0.005° , $N = 542$). Hence, flagella do not appear to play an important role for colony chirality. This is consistent with the results obtained for DH5 α , which is poorly motile, but shows significant chirality. In soft agar swimming assays, we also did not find any correlation between motility and chirality (data not shown). Supporting this observation is the fact that the highly chiral mutant, MG1655*fimflu*, showed lower motility than the parental strain MG1655 (data not shown).

In summary, the colony chirality significantly increased upon deletion of pili, colanic acid and curli fimbriae. In addition, it was enhanced upon double deletion of both pili and antigen 43. Hence, it appears that the more adhesive structures that are stripped off the cell, the more chiral the colony formation. Deletion of antigen 43 alone or of flagella lowered the chirality slightly, but not significantly, during colony expansion compared with the expansion of the mother strain. Supporting the small effect of flagella removal, chirality also is independent of swimming motility. We found a lower chirality of the strain MG1655*flufim* when grown at 30 $^\circ\text{C}$ (data not shown), and this may be linked to an increased expression of colanic acid and curli fimbriae at 30 $^\circ\text{C}$ (Prigent-Combaret *et al.*, 2000; Gualdi *et al.*, 2008).

Chirality and cell wall biosynthesis

During growth of *E. coli* an MreB guided insertion of material into the peptidoglycan has been suggested to cause a chiral ordering of the bacterial cell wall

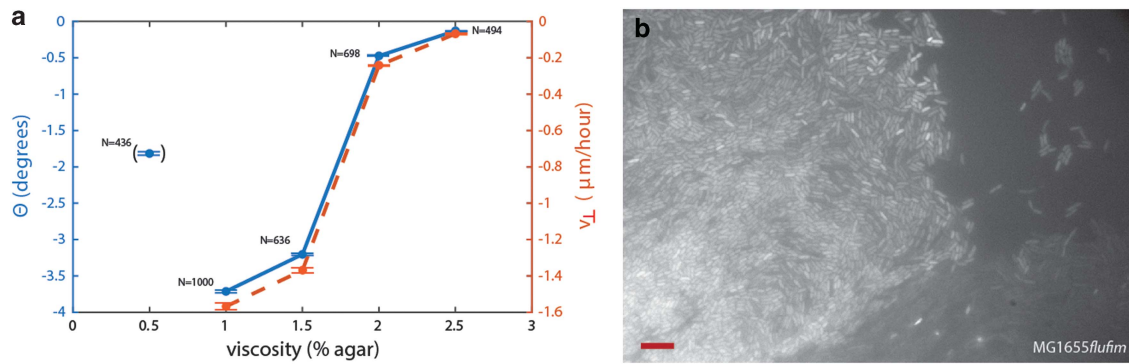


Figure 4 Effect of agar on chirality of sector boundaries and a microscopy image of the sector–sector and colony–agar boundaries. (a) Chirality measured θ (blue points and full line) and the border velocity, v_{\perp} (red points and punctuated line) are plotted for different agar concentrations; here MG1655flufim *E. coli* strain with high chirality was used. The number of boundaries analyzed, N , are noted for each experiment and the error bars signify one s.e.m. At 0.5 % agar, the colony growth was severely compromised compared with other experiments. To indicate this, we put the corresponding θ value in parenthesis. (b) Fluorescent microscopy image (the YFP channel) of sector–sector and colony–agar boundaries in the MG1655flufim strain. The scale bar is 10 μm .

and a left-handed twisting motion of the elongating *E. coli* cell (Wang *et al.*, 2012). This twisting motion of individual cells could affect the overall chirality of a colony. To investigate a possible relation between the twisting of bacteria during growth and chirality of the corresponding colony formation, we performed competition experiments with *B. subtilis* colonies. The cell wall structure of *B. subtilis* cells twist with an opposite handedness in comparison with *E. coli*; hence, we wanted to investigate whether this would cause the sector boundaries to twist in a direction opposite to that in *E. coli* colonies. Therefore, we performed competition experiments with the *B. subtilis* strains described in van Gestel *et al.* (2014). The sector boundaries between competing *B. subtilis* strains are slightly chiral returning a chirality angle $\theta = -1.1^{\circ}$ (s.e.m. = 0.03° , $N = 286$), an image from a *B. subtilis* competition experiment is shown in Supplementary Figure S2b. Hence, we did not see a correlation between the twisting motion of the underlying structure of the cell wall and the overall chirality during colony expansion; this could, however, be because the underlying cell wall structures were masked by extracellular constituents in *B. subtilis*. In addition, it should be noted that the *B. subtilis* grow with a morphology quite different from *E. coli* (see example in Supplementary Figure S2) which complicates a direct comparison. Interestingly, we did find that sector boundaries became coarser and less chiral when exopolysaccharides were expressed.

Effect of varying agar concentration

We also investigated the dependence of colony chirality on agar concentration. To this end MG1655flufim was plated on agar plates with agar concentrations in the interval 0.5–2.5%. The observed relation between θ and viscosity is shown in Figure 4a. For an agar concentration of 0.5% the sector boundaries became less clear and the colonies tended to spread into the agar (as opposed to

spreading on top of the agar); hence, the nature of colony growth differed significantly from growth at other concentrations. With this point excluded, we found that increasing the agar concentration decreased chirality and θ approached zero at the highest agar concentration tested. These results suggest that the interaction with the surface may become too strong to allow the emergence of chiral behavior. In addition, agar concentration may affect expression of surface structures and thereby influence colony chirality.

Colony chirality, θ , is a composite measure of chiral velocity tangential to the expansion front, v_{\perp} and the radial outward velocity of colony growth, v_{\parallel} ($v_{\perp}/v_{\parallel} = \tan(\theta)$). Since both velocities could depend on the agar concentration, we directly measured v_{\parallel} and then computed v_{\perp} from the measured values of θ . The radial velocity component, v_{\parallel} , increased by 25% ($24\text{--}30\ \mu\text{m h}^{-1}$), as the agar concentration increased from 1 to 2.5%. Simultaneously, the vertical velocity component, v_{\perp} , decreased by 96% from (-1.6 to $-0.07\ \mu\text{m h}^{-1}$). Hence, the change of v_{\perp} was substantially larger than the change of v_{\parallel} and it is clear from Figure 4a that v_{\perp} is proportional to θ . The radial colony growth is only slightly perturbed by changing the viscosity and diffusion of nutrients in accordance with mechanical predictions (Farrell *et al.*, 2013). However, the chirality is highly affected.

Microscopic alignment during colony expansion

Individual rod-shaped *E. coli* cells align themselves parallel to the boundaries between the two competing strains; see Figure 4b for an image for the MG1655flufim. This parallel alignment of mother and daughter cells can be explained by a mechanism where a large number of cells competing for space causes cells to push against each other, and this mechanical instability leads to buckling and folding of the parallel cells lines (Rudge *et al.*, 2012). Away from the boundaries, we often observed cell

alignment at the rim of the colony towards the empty agar plate (Figure 4b) as reported previously (Shapiro and Dworkin, 1988; Shapiro and Hsueh, 1989;

Shapiro, 1995). In contrast, colonies of the spherical *S. cerevisiae* (budding yeast) show neither appreciable alignment nor chirality (Hallatschek *et al.*, 2007;

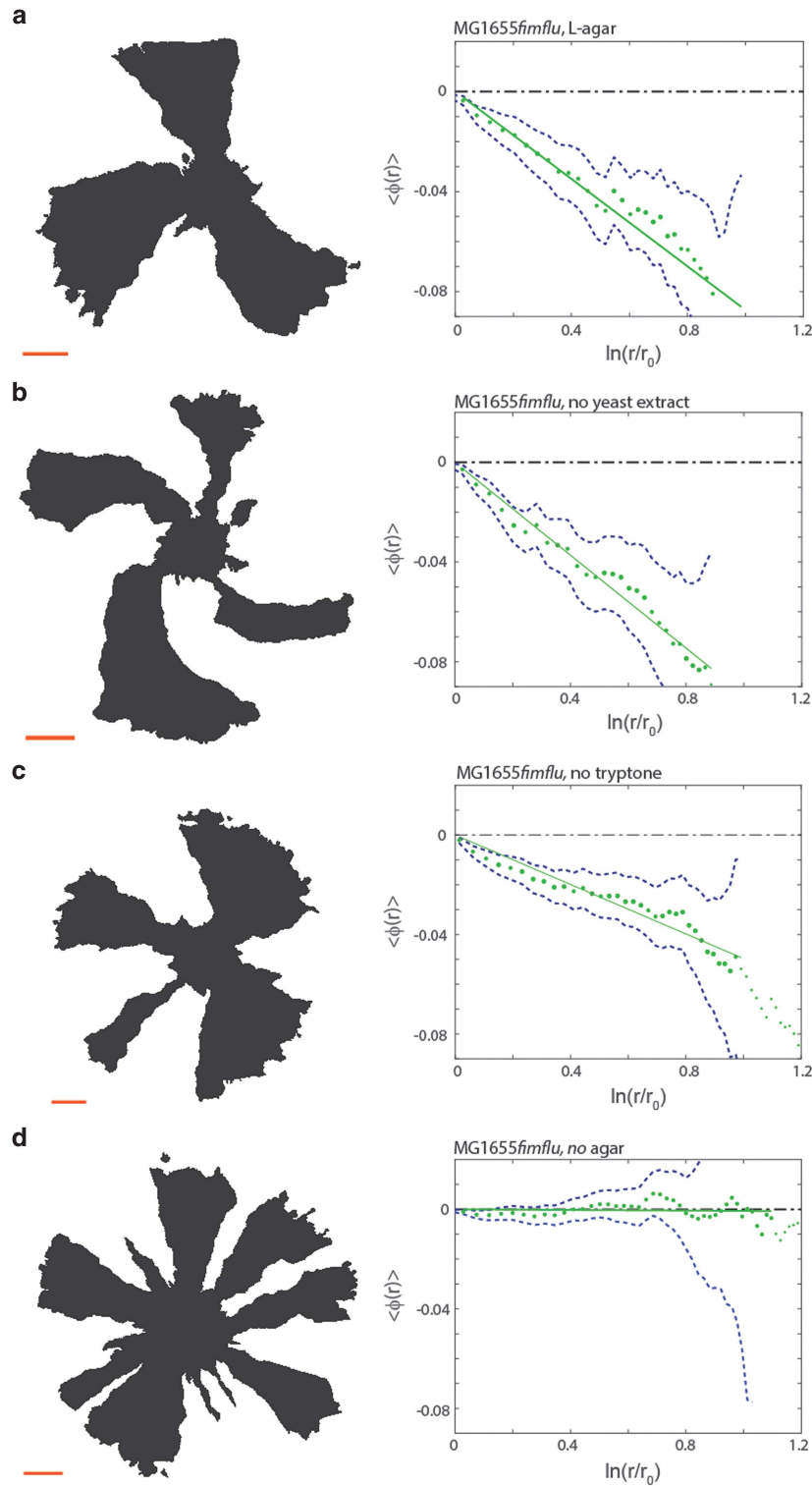


Figure 5 Effect of substratum substitution. The left panels show examples of intensity thresholded images (displayed from the agar side) of the *E. coli* MG1655*fimflu* mutant in competition experiments on LB medium with systematic replacements of constituents; the scale bars are 1 mm. The right panels show the average boundary angle, $\langle\phi(r)\rangle$ (green dotted line), a linear fit (full green line) and one s.e.m. (blue punctuated line). (a) Results from control competition experiments on normal LB medium (1.2% agar) ($N=222$). (b) Replacement of yeast extract by $1\ \mu\text{g ml}^{-1}$ B1, 1 mM MgCl_2 , and 10 mM sodium phosphate ($N=223$). (c) Replacement of tryptone and NaCl by 1% decolorized casamino acids ($N=337$). (d) Replacement of agar by 1.2% agarose ($N=410$).

Korolev *et al.*, 2012; Müller *et al.*, 2014).

To investigate whether macroscopic chirality is determined by the geometry of the microorganism, we performed chirality assays with *S. pombe* (fission yeast). *S. pombe* cells are rod shaped albeit larger than *E. coli* and with a smaller aspect ratio. The boundaries separating competing *S. pombe* strains are not chiral (see Supplementary Figure S2a); hence, the rod-like shape is not sufficient to cause chiral growth of microorganism colonies. An elongated shape could, however, still be necessary for asymmetric growth as shown, for example, for morphogenesis in *Drosophila* (Taniguchi *et al.*, 2011).

Trypsin-sensitivity assays

Having observed that systematic removal of features from the *E. coli* cell surface gave rise to increasingly chiral colony growth, we searched for the mediator of the chirality. To this end, we grew the double deletion strain, MG1655 *fimflu*, on agar plates with trypsin (0.025% w/v). Trypsin is a pancreatic serine protease that cleaves peptide chains on the carboxyl side of lysine and arginine and digests proteins on the cell surface. The overall patterning and the radial growth rate of the colony were unaffected by the presence of trypsin; however, it had a large effect on the chirality of the population boundaries: The chirality angle of the MG1655 *fimflu* strain was reduced to $\theta = -1.0^\circ$ (s.e.m. = 0.005° , $N = 1952$) (Supplementary Figure S3) similar to the parental level (MG1655). Hence, a non-essential protein component of the bacterial cell surface or the substratum mediates the chiral growth. To test whether trypsin enzymatically modulated the substratum, we let trypsin (0.025%) incubate with LB agar (1.2%) before pouring plates with or without the trypsin-inhibitor AEBSF (0.02%). We found that even in the presence of AEBSF chirality of the MG1655 *fimflu* strain decreased to a level below parental level (data not shown). This observation suggests trypsin enzymatically modulated the substratum, and maybe the cell surface, and indirectly modulates chirality.

Substratum substitution experiments

To identify which component of the substratum mediates chiral growth we performed a stepwise substitution of each LB agar constituent. For each substitution, we performed competition experiments using the MG1655 *fimflu* strain and the results are shown in Figure 5. The following experiments were performed: A control experiment, Figure 5a, on normal LB agar (1.2%) which gave $\theta = -4.0^\circ$ (s.e.m. = 0.05° , $N = 222$). Replacement of yeast extract by vitamin B₁, magnesium chloride and sodium phosphate (Figure 5b) gave $\theta = -6.7^\circ$ (s.e.m. = 0.07° , $N = 223$). Replacement of tryptone by decolorized casamino acids (Figure 5c) gave $\theta = -4.0^\circ$ (s.e.m. =

0.05° , $N = 337$), a chirality identical to the control. However, replacing 1.2% agar with 1.2% agarose (Figure 5d) reduced chirality dramatically to $\theta = -0.1^\circ$ (s.e.m. = 0.01° , $N = 410$), independent of viscosity of the plates (data not shown). Taken together, these results show that the substratum composition modulates chirality either directly, for example, via an intrinsic chiral structure, or indirectly, for example, by altering gene expression or by mediating adhesion of bacteria with accessible chiral surface structures. We cannot exclude the possibility that the change in composition also affects the metabolism/phenotype of the cells. It is well known that small changes in growth medium and solidifying agents can alter gene expression dramatically. This is also true for the expression of cell surface structures. Hence, the change in medium composition may also affect the expression of certain cell surface structures in the cells.

Discussion

Some *E. coli* strains display chiral borders between competing subpopulations during colony expansion. We found that a systematic removal of extracellular constituents involved in biofilm formation, cell–cell and cell–surface adhesion increased chirality. Deletion of pili, curli fimbriae and colanic acid in *E. coli* MG1655 produced sector boundaries that were more chiral. Loss of two extracellular constituents (antigen 43 and pili) increased chirality further. Interestingly, *E. coli* MG1655 colony formation displayed relatively low chirality compared with the other K-12 strain, DH5 α . This could be due to inadvertent selection in the construction of *E. coli* DH5 α , which is known to be a poor biofilm former and likely lacks certain extracellular structures present in MG1655. These results support a view that intimate contact with the substratum is responsible for the chirality in *E. coli*, and the adhesive surface structures are masking it by physical shielding or by reducing cell movement during colony growth.

Some of the largest surface structures on *E. coli* are the pili, which are thin, rigid and adhesive organelles. They are found on many *E. coli* strains and other members of the *Enterobacteriaceae* (Klemm and Krogfelt, 1994) and are important for adherence to mammalian host tissues (Krogfelt *et al.*, 1990; Connell *et al.*, 1996) and for biofilm formation (Pratt and Kolter, 1998; Schembri and Klemm, 2001). On a piliated cell, these peritrichous surface structures can radiate to a distance of up to 2 μm from the cell surface (Figure 3a), making them prime candidates for a physical masking of an underlying cell envelope structure. Accordingly, we found that the deletion of piliating genes in MG1655 (MG1655 *fim*) significantly increased colony chirality. Interestingly, pili expression, which is subject of phase variation, that is, individual cells can switch between a piliated and non-piliated state, is normally

considered low on solid media. Nevertheless, the presence of pili clearly decreases chirality.

Antigen 43, the product of the *flu* gene, is a self-recognizing auto-transporter protein. It confers auto-aggregation giving rise to a characteristic frizzy colony morphology on solid media as well as flocculation of cells in static liquid cultures (Hasman *et al.*, 1999). Antigen 43 is known to be important for bacterial biofilm formation (Danese *et al.*, 2000; Kjærgaard *et al.*, 2000; O'Toole *et al.*, 2000) and for cellular chain formation (Vejborg and Klemm, 2009). Interestingly, antigen 43 mediated auto-aggregation is inhibited by the presence of pili on the cells (Hasman *et al.*, 1999). Consistent with this finding, we observed little change of chirality in the MG1655/*flu* mutant, but a significant enhancement in θ of the double *flu*, *fim* mutant (MG1655/*flu-fim*). Overexpression of pili in the MG1655/*flufim* also could not fully reverse the phenotype, clearly suggesting that antigen 43 also plays an important role, and both surface structures mask the underlying chirality of the cell.

The deletion of curli fimbriae (MG1655/*csgAB*) or colanic acid (MG1655/*cps*) also made sector boundaries significantly more chiral. This increase in chirality was somewhat surprising, given that curli and colanic acid are not typically expressed in K-12 strains at 37 °C (Olsén *et al.*, 1989). Given that the expression of many surface structures is intricately connected, we cannot exclude the possibility that the deletion of these genes could affect the expression of other surface structures. It should also be noted that MG1655/*cps* was independently derived and could have acquired additional genetic and phenotypic variations, which might obscure direct comparison with our version of *E. coli* MG1655.

Flagella-dependent motility is known to play a major role in chiral spreading in other bacteria. For example, *Proteus mirabilis* produces chiral swarming patterns via this mechanism (Xue *et al.*, 2011). *E. coli* also use flagella for active swimming, which can produce chiral motion due to hydrodynamic interactions (DiLuzio *et al.*, 2005). In addition, flagella are involved in biofilm formation (Pratt and Kolter, 1998), and they may play a structural role in colony biofilms (Serra *et al.*, 2013). Not surprisingly, flagella have been put forward as the primary suspect for chiral behavior in bacteria. For instance, for *Paenibacillus dendritiformis*, it is proposed in Ben-Jacob *et al.* (1995) that chirality is caused by flagella and is mediated by strong cell–cell interaction. We found that chirality decreases slightly, and not statistically significantly, upon flagella removal. However, there was no link between the magnitude of chirality and flagella-mediated motility (as determined in swimming motility assays); in particular, the most chiral strains (DH5 α and MG1655/*flufim*) were poorly motile. Taken together, our data suggest that flagella, for these types of assays, contributes little to chirality in *E. coli* colonies under the studied growth conditions.

The macroscopic handedness of the chiral borders between competing *E. coli* strains is opposite of the microscopic handedness of the growth of the peptidoglycan layer (Wang *et al.*, 2012). Hence, although chirality is increased upon the removal of surface structures that possibly mask underlying cell wall structures, there is no direct link between the spiral growth direction of the peptidoglycan and the chirality of colony borders. This was confirmed by the same chirality direction of *E. coli* and *B. subtilis* despite the opposite directions of cell wall biosynthesis in these two species. However, spiral growth has in *Myxococcus xanthus* been linked to another kind of motility (gliding) than employed by *E. coli*.

Similar to chirality in regular matter (de Gennes and Prost, 1995), chirality in bacterial colonies may require parallel geometric ordering of cells during cell division and colony expansion (Shapiro and Hsueh, 1989; Su *et al.*, 2010), which is clearly visible at the border between competing strains (Figure 4b). The composition of the substratum has been shown to control cell morphogenesis (Su *et al.*, 2010). Hence, the alignment of mother–daughter cells is probably affected by the cells' ability to attach to the substratum and possibly to each other. Supporting the importance of the substratum for chiral growth, we also found that interchanging agar with agarose in the substratum reduced chirality. This suggests that the agar polymer network itself could be chiral or that it includes chiral entities. However, it could also be that cell–surface adhesion were modulated or that gene expression was changed.

Active chiral processes play an important role in pattern formation as well as during embryonic development (Coutelis *et al.*, 2014). While the ecological role of chirality in the microbial world is still poorly understood, chiral phenotypes could be functional or even advantageous to microbial communities as they are during embryogenesis. Indeed, one can view microbial colonies as a stepping-stone towards multi-cellularity (Shapiro and Dworkin, 1988), and, therefore, related pattern forming mechanisms could be required. Consistent with these ideas, a switch of chirality has been reported in bacteria as a response to changes in the environmental conditions (Ben-Jacob *et al.*, 2000) and during tumorigenesis (Wan *et al.*, 2011). We found that *E. coli* employs a mechanisms to generate chirality distinct from higher organisms, which often rely on the asymmetric orientation of the mitotic spindle during cell division (Grande and Patel, 2009) or, in the case of *Caenorhabditis elegans*, on chiral torque generation in the cortex (Naganathan *et al.*, 2014). Although the microscopic mechanisms for chirality in bacteria are distinct, some conclusions of our work could generalize to other settings. In particular, we established an experimental approach to examine the role of surface structure and adhesion in chiral pattern formation.

Conclusions

Chirality is a general property of biological systems which is shared by very different species as the Gram-positive *B. subtilis* and the Gram-negative *E. coli*. Efficient adaptation of bacterial colonies to changing environmental conditions requires cooperative behavior and self-organization, which is based on the exchange of information. The different communication channels range from direct physical and chemical interaction to indirect interactions through chemotaxis and trails on the agar. Pattern formation in microbial colonies is a way to exchange information between the microscopic level and the macroscopic level. In other words, an exchange between the individual cells and the colony. Therefore, the full understanding of chiral growth, its origins and implications must focus on the relation between the individual cell and the colony organization. For *E. coli* colonies expanding on a substratum, the border between competing strains often exhibits chirality and always in a left-handed manner (when observed from the agar side). We find a correlation between the expression of surface structures involved in cell–cell and cell–surface adhesion or biofilm formation and the macroscopic degree of chirality; the more surface structures expressed, the less chiral the border between competing strains. This result was obtained by comparing pattern formation of MG1655 to pattern formation of MG1655 mutants with deletion of pili, curli fimbriae, colanic acid and antigen 43. Furthermore, we found chirality to be independent of flagella-mediated motility. The DH5 α strain had more chiral borders than any of the MG1655 mutants, possibly because DH5 α is a poorer biofilm former. We also find that substituting agar with agarose reduces chirality, suggesting that chirality largely depend on substratum composition. The exact mechanism by which the substratum mediates the chiral growth will be a subject of future investigation. Overall, intimate cell–substratum adhesion is necessary for chirality and extracellular features, such as fimbriae or biofilm, weakens this contact. Our results link macroscopic chiral colony formation to the microscopic biological features of *E. coli*'s cell wall and the substratum and suggest new ecological roles for several membrane-associated proteins during colony growth.

Conflict of Interest

The authors declare no conflict of interest.

Acknowledgements

We thank G Thon and J van Gestel for providing us with *S. pombe* and *B. subtilis* strains, respectively. We also thank the anonymous referee for valuable comments. We are grateful to the Lundbeck foundation, the Danish National Research Foundation grant number DNR116,

the Carlsberg foundation and the Danish Research Councils for the financial support.

References

- Ali A, Grosskinsky S. (2010). Pattern formation through genetic drift at expanding population fronts. *Adv Complex Syst* **13**: 349–366.
- Bachmann BJ. (1996). Derivations and genotypes of some mutant derivatives of *Escherichia coli* K-12. In: Neidhardt FC editor, *Escherichia Coli and Salmonella Typhimurium: Cellular and Molecular Biology*. ASM Press: Washington, DC, pp 2460–2488.
- Beloin C, Michaelis K, Lindner K, Hacker J, Ghigo J, Dobrindt U *et al.* (2006). The transcriptional antiterminator RfaH represses biofilm formation in *Escherichia coli*. *J Bacteriol* **188**: 1316–1331.
- Ben-Jacob E, Cohen I, Levine H. (2000). Cooperative self-organization of microorganisms. *Adv Phys* **49**: 395–554.
- Ben-Jacob E, Cohen I, Shochet O, Tenenbaum A. (1995). Cooperative formation of chiral patterns during growth of bacterial colonies. *Phys Rev Lett* **75**: 2899–2902.
- Chang ACY, Cohen SN. (1978). Construction and characterization of amplifiable DNA cloning vectors derived from P15A cryptic plasmid. *J Bacteriol* **134**: 1141–1156.
- Connell I, Agace W, Klemm P, Schembri M, Märdil S, Svanborg C. (1996). Type 1 fimbrial expression enhances *Escherichia coli* virulence for the urinary tract. *Proc Natl Acad Sci USA* **93**: 9827–9832.
- Coutelis J-B, González-Morales N, Géminard C, Noselli S. (2014). Diversity and convergence in the mechanisms establishing L/R asymmetry in metazoa. *EMBO Rep* **15**: 926–937.
- Danese PN, Pratt LA, Dove SL, Kolter R. (2000). The outer membrane protein, Antigen 43, mediates cell-to-cell interactions within *Escherichia coli* biofilms. *Mol Microbiol* **37**: 424–432.
- Datsenko KA, Wanner BL. (2000). One-step inactivation of chromosomal genes in *Escherichia coli* K-12 using PCR products. *Proc Natl Acad Sci USA* **97**: 6640–6645.
- Diderichsen B. (1980). flu, A metastable gene controlling surface properties of *Escherichia coli*. *J Bacteriol* **141**: 858–867.
- DiLuzio WR, Turner L, Mayer M, Garstecki P, Weibel DB, Berg HC *et al.* (2005). *Escherichia coli* swim on the right-hand side. *Nature* **435**: 1271–1274.
- Edie ES. (1914). On the resistance of trypsin solutions to heat. *Biochem J* **8**: 84–89.
- Farrell FDC, Hallatschek O, Marenduzzo D, Waclaw B. (2013). Mechanically driven growth of quasi-two-dimensional microbial colonies. *Phys Rev Lett* **111**: 168101.
- Flack HD. (2009). Louis Pasteurs discovery of molecular chirality and spontaneous resolution in 1848, together with a complete review of his crystallographic and chemical work. *Acta Crystallogr Sect A Found Crystallogr* **65**: 371–389.
- Frank FC. (1953). On spontaneous asymmetric synthesis. *Biochim Biophys Acta* **11**: 459–463.

- Gayathri VS, Rao M. (2005). Fluctuation induced chiral symmetry breaking in autocatalytic reaction-diffusion systems. *Europhys Lett* **80**: 4.
- de Gennes P-G, Prost J. (1995). *The Physics of Liquid Crystals (International Series of Monographs on Physics)*. Oxford University Press: New York.
- van Gestel J, Weissing FJ, Kuipers OP, Kovács AT. (2014). Density of founder cells affects spatial pattern formation and cooperation in *Bacillus subtilis* biofilms. *ISME J* **8**: 2069–2079.
- Grande C, Patel NH. (2009). Nodal signalling is involved in left-right asymmetry in snails. *Nature* **457**: 1007–1011.
- Gualdi L, Tagliabue L, Bertagnoli S, Ieranò T, De Castro C, Landini P. (2008). Cellulose modulates biofilm formation by counteracting curli-mediated colonization of solid surfaces in *Escherichia coli*. *Microbiology* **154**: 2017–2024.
- Hallatschek O, Hersen P, Ramanathan S, Nelson DR. (2007). Genetic drift at expanding frontiers promotes gene segregation. *Proc Nat Acad Sci USA* **104**: 19926–19930.
- Hallatschek O, Nelson DR. (2010). Life at the front of an expanding population. *Evolution* **64**: 193–206.
- Hasman H, Chakraborty T, Klemm P. (1999). Antigen-43-mediated autoaggregation of *Escherichia coli* is blocked by fimbriation. *J Bacteriol* **181**: 4834–4841.
- Hasman H, Schembri MA, Klemm P. (2000). Antigen 43 and type 1 fimbriae determine colony morphology of *Escherichia coli* K-12. *J Bacteriol* **182**: 1089–1095.
- Huntley HE. (1970). *The Divine Proportion: A Study in Mathematical Beauty*. Courier Dover Publications: New York, NY, USA.
- Jafarpour F, Biancalani T, Goldenfeld N. (2015). Noise-induced mechanism for biological homochirality of early life self-replicators. *Phys Rev Lett* **115**: 1–5.
- Kjærgaard K, Schembri MA, Ramos C, Molin S, Klemm P. (2000). Antigen 43 facilitates formation of multispecies biofilms. *Environ Microbiol* **2**: 695–702.
- Klemm P, Krogfelt K. (1994). Type 1 fimbriae of *Escherichia coli*. In: Klemm P (ed). *Fimbriae: Adhesion, Genetics, Biogenesis and Vaccines*. CRC Press: Boca Raton, pp 9–26.
- Korolev K, Nelson D. (2011). Competition and cooperation in one-dimensional stepping-stone models. *Phys Rev Lett* **107**: 1–5.
- Korolev KS, Avlund M, Hallatschek O, Nelson DR. (2010). Genetic demixing and evolution in linear stepping stone models. *Rev Mod Phys* **82**: 1691–1718.
- Korolev KS, Müller MJI, Karahan N, Murray AW, Hallatschek O, Nelson DR et al. (2012). Selective sweeps in growing microbial colonies. *Phys Biol* **9**: 1–38.
- Korolev KS, Xavier JB, Nelson DR, Foster KR. (2011). A quantitative test of population genetics using spatiogenetic patterns in bacterial colonies. *Am Nat* **178**: 538–552.
- Kreft J-U. (2004). Conflicts of interest in biofilms. *Biofilms* **1**: 265–276.
- Krogfelt KA, Bergmans H, Klemm P. (1990). Direct evidence that the FimH protein is the mannose-specific adhesion of *Escherichia coli* type 1 fimbriae. *Infect Immun* **58**: 1995–1998.
- Kuroda R, Endo B, Abe M, Shimizu M. (2009). Chiral blastomere arrangement dictates zygotic left-right asymmetry pathway in snails. *Nature* **462**: 790–794.
- Mitri S, Clarke E, Foster KR. (2015). Resource limitation drives spatial organization in microbial groups. *ISME J* **10**: 1–12.
- Müller MJI, Neugeboren BI, Nelson DR, Murray AW. (2014). Genetic drift opposes mutualism during spatial population expansion. *Proc Natl Acad Sci USA* **111**: 1037–1042.
- Naganathan SR, Fürthauer S, Nishikawa M, Jülicher F, Grill SW. (2014). Active torque generation by the actomyosin cell cortex drives left–right symmetry breaking. *Elife* **3**: 1–16.
- O’Toole G, Kaplan HB, Kolter R. (2000). Biofilm formation as microbial development. *Annu Rev Microbiol* **54**: 49–79.
- Oh YJ, Hubauer-Brenner M, Gruber HJ, Cui Y, Traxler L, Siligan C et al. (2016). Curli mediate bacterial adhesion to fibronectin via tensile multiple bonds. *Sci Rep* **6**: 33909.
- Olsén A, Jonsson A, Normark S. (1989). Fibronectin binding mediated by a novel class of surface organelles on *Escherichia coli*. *Nature* **338**: 652–655.
- Owen P, Caffrey P, Josefsson LG. (1987). Identification and partial characterization of a novel bipartite protein antigen associated with the outer membrane of *Escherichia coli*. *J Bacteriol* **169**: 3770–3777.
- Pratt LA, Kolter R. (1998). Genetic analysis of *Escherichia coli* biofilm formation: roles of flagella, motility, chemotaxis and type I pili. *Mol Microbiol* **30**: 285–293.
- Prigent-Combaret C, Prensier G, Le Thi TT, Vidal O, Lejeune P, Dorel C. (2000). Developmental pathway for biofilm formation in curli-producing *Escherichia coli* strains: role of flagella, curli and colanic acid. *Environ Microbiol* **2**: 450–464.
- Prigent-Combaret C, Vidal O, Dorel C, Lejeune P. (1999). Abiotic surface sensing and biofilm-dependent regulation of gene expression in *Escherichia coli*. *J Bacteriol* **181**: 5993–6002.
- Reisner A, Haagenen JA, Schembri MA, Zechner EL, Molin S. (2003). Development and maturation of *Escherichia coli* K-12 biofilms. *Mol Microbiol* **48**: 933–946.
- Rudge TJ, Steiner PJ, Phillips A, Haseloff J. (2012). Computational modeling of synthetic microbial biofilms. *ACS Synth Biol* **1**: 345–352.
- Saito Y, Sugimori T, Hyuga H. (2007). Stochastic approach to enantiomeric excess amplification and chiral symmetry breaking. *J Phys Soc Japan* **76**: 1–14.
- Saldaña Z, Xicohtencatl-Cortes J, Avelino F, Phillips AD, Kaper JB, Puente JL et al. (2009). Synergistic role of curli and cellulose in cell adherence and biofilm formation of attaching and effacing *Escherichia coli* and identification of Fis as a negative regulator of curli. *Env Microbiol* **11**: 992–1006.
- Schembri MA, Klemm P. (2001). Coordinate gene regulation by fimbriae-induced signal transduction. *EMBO J* **20**: 3074–3081.
- Serra DO, Richter AM, Klauck G, Mika F, Hengge R. (2013). Microanatomy at cellular resolution and spatial order of physiological differentiation in a bacterial biofilm. *MBio* **4**: 1–12.
- Shapiro JA. (1995). The significance of bacterial colony patterns. *BioEssays* **17**: 597–607.
- Shapiro JA, Dworkin M. (1988). Bacteria as multicellular organisms. *Sci Am* **6**: 82–89.

- Shapiro JA, Hsut C. (1989). Escherichia coli K-12 cell-cell interactions seen by time-lapse video. *J Bacteriol* **171**: 5963–5974.
- Su P-T, Yen P-W, Wang S-H, Lin C-H, Chiou A, Syu W-J. (2010). Factors affecting daughter cells' arrangement during the early bacterial divisions. *PLoS One* **5**: e9147.
- Taniguchi K, Maeda R, Ando T, Okumura T, Nakazawa N, Hatori R *et al.* (2011). Chirality in planar cell shape. *Science (80-)* **339**: 339–342.
- Vejborg RM, Klemm P. (2009). Cellular chain formation in Escherichia coli biofilms. *Microbiology* **155**: 1407–1417.
- Vidal O, Longin R, Prigent-Combaret C, Dorel C, Hooreman M, Lejeune P. (1998). Isolation of an Escherichia coli K-12 mutant strain able to form biofilms on inert surfaces: Involvement of a new ompR allele that increases curli expression. *J Bacteriol* **180**: 2442–2449.
- Wan LQ, Ronaldson K, Park M, Taylor G, Zhang Y, Gimble JM *et al.* (2011). Micropatterned mammalian cells exhibit phenotype-specific left-right asymmetry. *Proc Natl Acad Sci USA* **108**: 12295–12300.
- Wang S, Furchtgott L, Huang KC, Shaevitz JW. (2012). Helical insertion of peptidoglycan produces chiral ordering of the bacterial cell wall. *Proc Natl Acad Sci USA* **109**: E595–E604.

- Wolfe AJ, Berg HC. (1989). Migration of bacteria in semisolid agar. *Proc Natl Acad Sci USA* **86**: 6973–6977.
- Xue C, Budrene EO, Othmer HG. (2011). Radial and spiral stream formation in proteus mirabilis colonies. *PLoS Comput Biol* **7**: 22–26.
- Yu D, Ellis HM, Lee E-C, Jenkins Na, Copeland NG, Court DL. (2000). An efficient recombination system for chromosome engineering in Escherichia coli. *Proc Natl Acad Sci USA* **97**: 5978–5983.



This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivs 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by-nc-nd/4.0/>

© The Author(s) 2017

Supplementary Information accompanies this paper on The ISME Journal website (<http://www.nature.com/ismej>)